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Substitute for form 1449A/PTO  <h2 style="text-align: center; margin: 0;">INFORMATION DISCLOSURE STATEMENT BY APPLICANT</h2> <p style="text-align: center; margin: 0;"><i>(use as many sheets as necessary)</i></p>				<b>Complete if Known</b>	
Application Number		09/776,936			
Filing Date		December 22, 1998			
First Named Inventor		Scott Miller			
Group Art Unit		1621			
Examiner Name		Kumar, Shailendra			
Attorney Docket Number		BAYER-0006-P01			
Sheet	1	of	7		

U.S. PATENT DOCUMENTS					
Examiner Initials *	Cite No. <sup>1</sup>	U.S. Patent Document		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY
		Number	Kind Code <sup>2</sup> (if known)		
	A1	4,546,191		Nishiyama et al.	10-08-1985
	A2	6,380,218		Marafat et al.	04-03-2002
	A3	6,525,046		Cirillo et al.	02-25-2003
	A4	6,500,863		Jin et al.	12-31-2002
	A5	6,040,339		Yoshida et al.	03-21-2000
	A6	6,150,415		Hammock et al.	11-21-2000
	A7	6,178,399		Takebayashi et al.	01-23-2001
	A8	6,187,799		Wood et al.	02-13-2001
	A9	7,625,915		Dumas et al.	12-01-2002

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		Office <sup>3</sup>	Number <sup>4</sup>	Kind Code <sup>5</sup> (if known)				
	B1	DE	3529247	A1	Bayer AG	11-20-1986		A9
	B2	WO	90/02112		The Nutrasweet Company	03-08-1990		
	B3	EP	0690344	A1	Konica Corporation	01-03-1996		
	B4	WO	97/09973	A1	The Regents Of The University Of California	3-20-1997		
	B5	WO	98/20868	A1	The Picower Institute of Medical Research	05-22-1998		
	B6	WO	98/45268		Pfizer Products, Inc.	10-15-1998		
	B7	WO	99/28305	A1	E. I. Du Pont de Nemours & Co.	06-10-1999		
	B8	WO	00/43366	A1	Kirin Beer Kabushiki Kaisha	07-27-2000		A9
	B9	WO	00/56331	A1	Vertex Pharmaceuticals, Inc.	09-28-2000		
	B10	WO	02/14331		The University of Kansas	02-21-2002		
	B11	WO	02/062763	A2	Bayer Corporation	8-15-2002		

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Sheet 2 of 7

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	B12	WO	02/083628	A1	Boehringer Ingelheim Pharmaceuticals Inc.	10-24-2002		
	B13	WO	02/085857	A2	Bayer Corporation	10-31-2002		
	B14	WO	02/085859	A1	Bayer Corporation	10-31-2002		
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	B16	WO	03/099771	A2	Novartis AG	12-04-2003		

### NON PATENT LITERATURE DOCUMENTS

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	C1	ADJEL et al., "A phase I study of BAY 43-9006 and gefitinib in patients with refractory or recurrent non-small-cell lung cancer (NSCLC)," Meeting: 2005 ASCO Annual Meeting, Category: Developmental Therapeutics; Molecular Therapeutics, Subcategory: Antiangiogenic or Antimetastatic agents, Abstract No. 4510	
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	C9	CHANG et al., "BAY 43-9006 (Sorafenib) inhibitors ectopic (s.c.) and orthotopic growth of a murine model of renal adenocarcinoma (Renca) predominantly through inhibition of tumor angiogenesis," 96 <sup>th</sup> Annual Meeting, April 16-20, 2005, Anaheim/Orange County, CA	

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	C10	CLARK et al., "Safety and pharmacokinetics of the dual action raf kinase and vascular endothelial growth factor receptor inhibitor, BAY 43-9006, in patients with advanced, refractory solid tumors," <i>Clin. Cancer Res.</i> , 2005:11(15), 1 August 2005, 5472-5480	
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	C23	HEIM et al., "Antitumor effect and potentiation or reduction in cytotoxic drug activity in human colon carcinoma cells by the Raf kinase inhibitor (RKI) BAY 43-9006," <i>International Journal of Clinical Pharmacology and Therapeutics</i> , Vol. 41, No. 12/2003 (616-617)	
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	C40	WILHELM, S. M. et al., "A Novel Diphenylurea Raf-1 Kinase Inhibitor (RKI) Blocks the Raf/Mek/Erk Pathway in Tumor Cells," Proceedings of the American Association for Cancer Research, March 2001, Vol. 42	
	C41	REDMAN, A. M. ET AL., "P38 Kinase Inhibitors for the Treatment of Arthritis and Osteoporosis: Thienyl, Furyl and Pyrrolyl Ureas" <i>Bioorganic Medical Chemistry Letters</i> , 2001, Vol. 11 No. 1, pp. 9	
	C42	REGAN et al., "Pyrazole urea-based inhibitors of P38 MAP kinase: from lead compound to clinical candidate," <i>J. Med. Chem.</i> 45:2994-3008, 2002	
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	C45	RIEDL et al., "Potent Raf Kinase Inhibitors from the Diphenylurea Class: Structure Activity Relationships," 2001, <i>Proc. Amer. Assoc. Can. Res.</i> , Vol. 42 No. 923	
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	C53	THOMPSON ET AL., "Recent progress in targeting the Raf/MEK/ERK pathway with inhibitors in cancer drug discovery," <i>Current Opinion Pharmacology</i> , Aug 2005, Vol. 5, No. 4, pp. 350-356	
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	C58	Wisner et al., "Analogues of platelet activating factor. 7. Bis-aryl amide and bis-aryl urea receptor antagonists of PAF," <i>J. Med. Chem.</i> , 1992, 35, 4779-4789	
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Sheet 7 of 7

### Complete if Known

Application Number	09/776,936
Filing Date	December 22, 1998
First Named Inventor	Scott Miller
Group Art Unit	1621
Examiner Name	Kumar, Shailendra
Attorney Docket Number	BAYER-0006-P01

### NON PATENT LITERATURE DOCUMENTS

Examiner Initials *	Cite No. <sup>1</sup>	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T <sup>2</sup>
	C62	CUNNINGHAM, C. C. ET AL., "A Phase I Trial of H-ras Antisense Oligonucleotide ISIS 2503 Administered as a Continuous Intravenous Infusion in Patients with Advanced Carcinoma", 2001 American Cancer Society, Volume 92, Number 5, pages 1265-1271.	
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